

IN THE CLAIMS:

Please enter the following amended claims:

- C | 1. (Currently amended) A substantially Substantially-purified form of a the polypeptide, said polypeptide that comprising the amino acid amino-acid-sequence shown in SEQ ID NO. 13 or 14, a homologue thereof, a fragment thereof or a homologue of the fragment.
- C | 2. (Currently amended) The A-polypeptide according to claim 1, said polypeptide comprising that consists (comprising) of the amino acid amino-acid-sequence shown in SEQ ID NO. 13 or 14.
3. (Original) A cDNA encoding the polypeptide according to claim 1.
4. (Currently amended) The A-cDNA according to claim 3, said cDNA that comprising the nucleotide sequence shown in SEQ ID NO. 11 or 15, or a fragment cDNA that selectively hybridizes hybridized to said the cDNA.
- C 2 5. (Currently amended) The A-cDNA according to claim 3, said cDNA that comprising the nucleotide sequence shown in SEQ ID NO. 12, or a fragment cDNA that selectively hybridizes hybridized to said the cDNA.
6. (Currently amended) A replication or expression vector carrying the cDNA according to any one of claims claim-3 to 5.
7. (Original) A host cell transformed with the replication or expression vector according to claim 6.
8. (Currently amended) A method for producing a the polypeptide, comprising according to claim 1 or 2 which comprises culturing a host cell according to claim 7 under a

condition effective to express a the polypeptide encoded by the cDNA according to claim 1 or 2,
and collecting the polypeptide encoded by the cDNA.

9. (Original) A monoclonal or polyclonal antibody against the polypeptide according to claim 1 or 2.

10. (Currently amended) A pharmaceutical composition comprising a containing the polypeptide according to claim 1 or 2 ~~or the antibody according to claim 9~~, in association with a pharmaceutically acceptable diluent and/or carrier.

11. (Currently amended) A pharmaceutical composition for the treatment of abnormal growth of a smooth muscle cell, comprising containing a polypeptide according to claim 1 or 2, in association with a pharmaceutically acceptable diluent and/or carrier.

C1 12. (Currently amended) A pharmaceutical composition for the treatment of arteriosclerosis, restenosis after PTCA or myosarcoma, comprising a containing the polypeptide according to claim 1 or 2, in association with a pharmaceutically acceptable diluent and/or carrier.

13. (Currently amended) A screening method for an antagonist or agonist of the polypeptide according to claim 1 or 2, comprising

- a) contacting a cell with -with using the said polypeptide and a test compound,
- b) determining a result on cell growth of said contact, and
- c) comparing said result with a result from a control experiment where a cell is contacted with the polypeptide in the absence of the test compound.

14. (Previously added) A method for promoting smooth muscle growth or vasculogenesis which comprises administering to a mammalian subject a therapeutically effective amount of an antibody of claim 9.

C1 15. (Previously added) A pharmaceutical composition for promoting smooth muscle cell growth or vasculogenesis which comprises administering to a mammalian subject a therapeutically effective amount of an antibody of claim 9.